

METHOXYAMINE COMBINATIONS IN THE TREATMENT OF CANCER

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5 government has certain rights in the invention.

FIELD OF THE INVENTION

This invention generally relates to novel compositions and methods for the
treatment of certain cancers. Additionally, this invention relates to novel compositions
and methods to screen drugs useful for the treatment of certain cancers.

BACKGROUND

Cancer is a worldwide problem. The American Cancer Society estimates that
over one half million people will die from cancer in the United States alone in 1999.
As such, finding novel compositions and methods for the treatment of cancer is of
vital interest. The treatment of cancer falls into three general categories:
15 chemotherapy, radiation therapy and surgery. Often, therapies are combined since a
combination of therapies often increases the probability the cancer will be eradicated
as compared to treatment strategies utilizing a single therapy. Most typically, the
surgical excision of large tumor masses is followed by chemotherapy and/or radiation
therapy.

20 Chemotherapeutic agents can work in a number of ways. For example,
chemotherapeutic can work by interfering with cell cycle progression or by generating
DNA strand breaks. If the cancer cell is not able to overcome the cell cycle blockage
or cell injury caused by the therapeutic compound, the cell will often die via apoptotic
mechanisms. The use of a single chemotherapeutic agent in the treatment of cancer,
25 with or without surgery or radiation, has several disadvantages. First, the cells may
develop resistance to the chemotherapeutic agent. Such resistance results either in the
requirement for higher dosages of the drug and/or the renewed spread of the cancer.

Chemotherapeutic agents can be toxic to the patient. Therefore, there is a practical upper limit to the amount that a patient can receive. However, if two chemotherapeutic agents are used in concert, the dosage of any single drug can be lowered. This is beneficial to the patient since using lower levels of chemotherapeutic agents is generally safer for the patient. Additionally, cancer cells are less likely to generate resistance to the combination of drugs as they are to a single drug.

The design of drug combinations for the chemotherapeutic treatment of cancer should be approached with the goals of 1) finding a combination that is synergistic with and not merely additive to the first compound with respect to the elimination of the tumor, and 2) finding a second drug that does not potentiate the toxic effects of the first chemotherapeutic agent. These conditions require a great deal of empirical testing of agents known to have anticancer properties with agents that either may have anticancer properties, or that may augment the first agent in other ways. TMZ is currently employed in chemotherapeutic treatment of certain tumors. It works by dramatically increasing the mutation rate of cells undergoing DNA replication. Such cells, because of the high number of mutations which they have acquired as a result of the treatment with TMZ, are rapidly removed by apoptosis, thereby potentially eliminating the tumor. Some tumor cells are resistant to treatment by TMZ due to deficiencies in the mismatch repair (MMR) system in the cell. A defective MMR system prevents the cell from recognizing O⁶mG DNA adducts thereby making the cell resistant to elimination.

Baer *et al.*, in U.S. Pat.No. 5,731,304, note that the toxicity of temozolomide can be potentiated by agents that inhibit the enzyme O⁶-alkylguanine DNA alkyltransferase (ATase). In particular, they note that O⁶-benzylguanine (BG) can enhance the toxicity of temozolomide in certain cell lines that exhibit high levels of ATase (*e.g.* 300-fold in MAWI cells). However, in other cell lines that exhibit lower levels of ATase (*e.g.* U373 cells) the effect is significantly less.

Mitchell and Dolan (*Cancer Chemother Pharmacol* 32:59-63, 1993) note that temozolomide (TMZ) and an analogue, 5-(dimethyltriazeno)imidazole-4-carboxamide (DITC), can be effective in enhancing the anti-tumor effects of 1,3-bis(2-chloroethyl)-

1-nitrosourea (BCNU). TMZ and DITC work by depleting cells or tumors of O⁶-alkylguanine-DNA alkyltransferase (AGT). AGT is a DNA repair protein that selectively removes adducts from the O⁶ position of guanine in DNA by a stoichiometric transfer of the alkyl group to a cysteine moiety. Removal of the alkyl group from the DNA by methylation of the O⁶ position of the guanine effectively inactivates the AGT. As with the patent referred to above, the disclosed method is limited to specific cells or cancers.

Therefore, what is need is the development of novel therapies that utilize the synergistic properties of two or more therapeutic agents for the treatment of cancer that have a broader range of targets or a different range of targets than those combination therapies already known.

SUMMARY OF THE INVENTION

The invention contemplates that temozolomide (TMZ) and methoxyamine (MX) shall be used as a treatment for certain tumors that are resistant to treatment by TMZ alone. Additionally, the invention contemplates the TMZ and inhibitors of poly (ADP-ribose polymerase (PARP) shall be used as a treatment for certain tumors that are resistant to treatment by TMZ alone. The present invention generally comprises novel compositions to 1) screen for compounds that can potentiate or modulate the therapeutic effect of temozolomide (TMZ) alone or combined with methoxyamine (MX), 2) provide model systems for the study of cancer treatments by agents that modulate DNA repair mechanisms and 3) provide treatments for certain cancers.

With regard to the treatment of cancer, the present invention contemplates methods of treating cancer that utilize TMZ in conjunction with another agent that is capable of potentiating the toxic effect of the TMZ. More specifically, the invention relates to potentiating the effect of TMZ with agents that modify DNA adducts created by TMZ. Even more specifically, the invention relates to potentiating the effect of TMZ with agents that modify N⁷-methylguanine (N⁷mG) and N³-methyladenine (N³mA) DNA adducts. One example of such an agent is MX. The present invention is not limited by the order in which the agents are administered. In one embodiment,

the agents are administered sequentially. In another embodiment, the agents are administered as a combined formulation (*i.e.*, a formulation comprising methoxyamine and temezolomide). In one embodiment, the present invention contemplates a method of treating cancer comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising methoxyamine and iii) a second formulation comprising temozolomide; b) administering said first formulation to said patient; and c) administering said second formulation to said patient.

Additionally, the invention relates to potentiating temozolomide with agents that interfere with the ability of PARP to be effective in DNA repair by either competing with PARP, inhibiting PARP or degrading PARP. The present invention is not limited to the agent used to inhibit PARP activity. In one embodiment, PD128763 is used. In another embodiment 6-AN is used. In another embodiment, 3-AB is used. The present invention is not limited by the order in which the agents are administered. In one embodiment the agents are administered sequentially. In another embodiment, the agents are administered as a combined formulation. In one embodiment, the present invention contemplates a method of treating cancer comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising a poly-(ADP-ribose)-polymerase inhibitor and iii) a second formulation comprising temozolomide; b) administering said first formulation to said patient; and c) administering said second formulation to said patient.

The present invention is not limited to the method of administration of the treatment. In one embodiment, the treatment is administered orally. In another embodiment the treatment is administered intravenously. In yet another embodiment, the treatment is administered intraperitoneally. In yet another embodiment, the treatment is administered directly to the tumor by injection or, in the case of skin tumors, for example, by direct application of creams or ointments.

As noted above, the present invention also contemplates screening assays to identify drugs that augment the ability of temozolomide to inhibit tumor growth, drugs that augment the ability of temozolomide with methoxyamine to inhibit tumor growth, and drugs that augment the ability of temozolomide with PARP inhibitors to inhibit

tumor growth. A variety of assay formats are contemplated for testing the potential of compounds suspected of augmenting the anti-tumor effect of TMZ. In one embodiment, cells are pretreated with the compound suspected of augmenting the anti-tumor effect of temozolomide, followed by treatment with temozolomide. Cell growth and/or cell death are then measured to determine if there is an anti-tumor effect. In another embodiment, cells are treated with the compound suspected of augmenting the anti-tumor effect of temozolomide at approximately the same time as they are treated with temozolomide. Cell growth and/or cell death are then measured to determine if the regime had an anti-tumor effect. The invention is not limited to any particular measurement technique for apoptosis or cell growth. Various methods are envisioned. For example, mitosis can be measured by use of fluorescent dyes that intercalate into DNA, by the measurement of H³-thymidine incorporation or by colormetric assays. Such assays permit the use of high throughput screening methods. Apoptosis can be measured by the use of fluorescent dyes that intercalate into DNA, annexin-V staining of phosphatidyl serine residues of the cell surface or morphological changes in cell appearance.

The present invention contemplates a method of screening for compounds that augment the anti-tumor effect of temozolomide comprising: a) providing i) tumor cells, ii) a first formulation comprising a compound suspected of potentiating temozolomide and iii) a second formulation comprising temozolomide; b) contacting said cells with said first formulation; c) contacting said cells with said second formulation; and d) measuring the growth and death rates of said cells.

The present invention contemplates a method of screening for compounds that augment the anti-tumor effect of temozolomide comprising: a) providing i) an experimental animal and ii) tumor cells; iii) a first formulation comprising a compound suspected of potentiating temozolomide and iv) a second formulation comprising temozolomide; b) contacting said tumor cells with said experimental animal in such a way so that the tumor cells will grow in said animal so to make an animal with cancer; c) administering said first formulation to said animal with cancer; d)

administering said second formulation to said animal with cancer; and e) measuring the growth and death rates of said cancer cells.

It is not intended that the present invention be limited by the nature of the compounds to be screened in the screening assay of the present invention. For example, a variety of compounds including peptides, organic compounds and nonorganic compounds, are contemplated. Additionally, combinations of compounds are contemplated by the present invention.

In some embodiments, a method is provided comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising methoxyamine and iii) a second formulation comprising 1,3-bis (chloroethyl) 2-nitrosourea (BCNU); b) administering said first formulation to said patient; and c) administering said second formulation to said patient; wherein said methoxyamine is administered in an amount sufficient to potentiate toxicity of said BCNU. In some embodiments, said methoxyamine and said BCNU are administered sequentially, while in other embodiments said methoxyamine and said BCNU are administered as a formulation. In some embodiments, said methoxyamine and said BCNU are administered orally, while in other embodiments, said methoxyamine and said BCNU are administered intravenously.

In yet other embodiments, a formulation comprising methoxyamine and BCNU is contemplated.

In other embodiments, a method is provided comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising methoxyamine and iii) a second formulation comprising an anticancer drug or agent that exerts cytotoxicity mediated by oxidative DNA damage; b) administering said first formulation to said patient; and c) administering said second formulation to said patient; wherein said methoxyamine is administered in an amount sufficient to potentiate toxicity of said anticancer agent or drug. In some embodiments, said anticancer drug or agent is selected from the group consisting of bleomycin and adriamycin. In some embodiments, said methoxyamine and said anticancer drug or agent are administered

sequentially. In other embodiments, said methoxyamine and said anticancer drug or agent are administered as a formulation.

In other embodiments, a method is provided comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising methoxyamine and iii) a second formulation comprising an anticancer drug or agent selected from the group consisting of hypoxanthine, 5-FU, uracil, IUdR, bleomycin and adriamycin; b) administering said first formulation to said patient; and c) administering said second formulation to said patient; wherein said methoxyamine is administered in an amount sufficient to potentiate toxicity of said anticancer drug or agent. In some embodiments, said methoxyamine and said anticancer drug or agent are administered sequentially, while in other embodiments, said methoxyamine and said anticancer drug or agent are administered as a formulation. In some embodiments, said administration is oral administration, in other embodiments, said administration is intravenous administration.

In some embodiments, a formulation comprising methoxyamine and an anticancer drug or agent selected from the group consisting of hypoxanthine, 5-FU, uracil, IUdR, bleomycin and adriamycin is contemplated. In some embodiments, said anticancer drug or agent is IUdR. In some embodiments, a formulation comprising methoxyamine and hydrogen peroxide is contemplated.

In some embodiments, a method is provided comprising: a) providing i) a patient diagnosed with cancer and ii) a formulation comprising methoxyamine; b) administering said formulation to said patient; and c) treating said patient with gamma radiation; wherein said methoxyamine is administered in an amount sufficient to potentiate the toxicity of said gamma radiation. In some embodiments, said methoxyamine administration and gamma radiation treatment occur sequentially, while in other embodiments, said methoxyamine administration occurs essentially simultaneously with said radiation treatment.

In some embodiments, a method is provided comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising methoxyamine and iii) a second formulation comprising hydrogen peroxide; b) administering said first formulation to said patient; and c) administering said second formulation to said

patient; wherein said methoxyamine is administered in an amount sufficient to potentiate the cytotoxic effects of said hydrogen peroxide.

In other embodiments, a method is provided comprising: a) providing i) a patient diagnosed with cancer, ii) a first formulation comprising methoxyamine and iii) a second formulation comprising iododeoxyuridine (IUdR); b) administering said first formulation to said patient; and c) administering said second formulation to said patient; wherein said methoxyamine is administered in an amount sufficient to further increase the radiosensitivity of the tumor cells in said patient. In some embodiments, the method further comprises the step of d) treating said patient with radiation therapy. In some embodiments, said methoxyamine and said IUdR are administered sequentially, while in other embodiments, said methoxyamine and said IUdR are administered as a formulation. In some embodiments, said methoxyamine and said IUdR are administered orally, while in other embodiments, said methoxyamine and said IUdR are administered intravenously.

DESCRIPTION OF THE FIGURES

Figure 1. MX increases the sensitivity of colon cancer cells to TMZ. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 6 mM MX or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. ^{Fig 1A} A. HTC116 cells, ^{Fig 1B} B. SW480 cells. (■), TMZ alone; (♦), MX plus TMZ; (●), BG plus TMZ; (▲), BG and MX plus TMZ. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 2. Cytotoxicity of MMS enhanced by MX in colon cancer cell lines. Cells were treated with 0-3 mM MMS for 1 h or MMS and 6 mM MX plus 10 μ M BG for 2 h prior to 2 h exposure to MMS. ^{Fig 2A} A. HCT116 cells, ^{Fig 2B} B. SW480 cells. (■), MMS alone; (♦), MX plus MMS; (●), BG plus MMS; (▲), BG and MX plus MMS. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 3. Inhibitors of PARP enhance cytotoxicity of TMZ in MMR wt SW480 cells. ^{Fig 3A} A. Cells were treated with 0-1500 TMZ for 2 h or TMZ and 100 μ M PD128763 or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ

alone; (◆), BG plus TMZ; (●), PD128763 plus TMZ; (▲), BG and TMZ plus

PD128763. ^{Fig 3B} B. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 10 μ M 6-AN (pretreated for 48 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ.

(■), TMZ alone; (◆), BG plus TMZ; (●), 6-AN plus TMZ; (▲), BG and TMZ plus 6-AN. ^{Fig 3C} C. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 3 mM 3-AB

(pretreated for 24 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (◆), BG plus TMZ; (●), 3-AB plus TMZ; (▲), BG and TMZ plus 3-AB.

Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 4. Inhibitors of PARP enhance cytotoxicity of TMZ in MMR deficient HCT116 cells. ^{Fig 4A} A. Cells were treated with 0-1500 TMZ for 2 h or TMZ and 100 μ M PD128763 or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ

alone; (◆), BG plus TMZ; (●), PD128763 plus TMZ; (▲), BG and TMZ plus PD128763. ^{Fig 4B} B. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 100 μ M 6-AN (pretreated for 48 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ.

(■), TMZ alone; (◆), BG plus TMZ; (●), 6-AN plus TMZ; (▲), BG and TMZ plus 6-AN. ^{Fig 4C} C. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 3 mM 3-AB (pretreated for 24 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■),

TMZ alone; (◆), BG plus TMZ; (●), 3-AB plus TMZ; (▲), BG and TMZ plus 3-AB.

Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 5. Synergy analysis of the interaction between TMZ and MX ^(Fig 5A) (A) or PD128763 ^(Fig 5B) (B) in HTC116 cells.

Figure 6. Survival fraction of human colon cancer cell lines after exposure to

BCNU plus MX. Cells were treated with 0-100 μ M BCNU for 2 h or BCNU and 6 mM MX plus 10 μ M BG for 2 h prior to 2 h exposure to BCNU. ^{Fig 6A} A. HCT116 cells;

^{Fig 6B} B. SW480 cells. (■), BCNU alone; (◆), MX plus BCNU; (●), BG plus BCNU; (▲), BG plus MX plus BCNU. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 7. Survival fraction of human colon cancer cell lines after exposure to BCNU plus PD128763. Cells were treated with 0-100 μ M BCNU for 2 h or BCNU and 100 μ M PD 128763 plus 10 μ M BG for 2 h prior to 2 h exposure to BCNU. HCT116 cells; B. SW480 cells. (■), BCNU alone; (◆), PD128763 plus BCNU; (●), BG plus BCNU; (▲), BG and PD128763 plus BCNU; (▶), BG, PD128763 and MX plus BCNU. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 8. Distribution of cell cycle and apoptosis in colon cancer cell lines after treatment with TMZ plus modifiers. A. MMR wt SW480 cells; B. MMR deficient HCT116 cells.

Figure 9. PARP cleavage in colon cancer cells treated with TMZ plus modifiers. A. Controls, as labeled. B. SW480 cells; C. HCT116 cells; Lane 1, untreated; 2, TMZ plus MX; 3, TMZ plus MX plus BG; 4, TMZ plus 3-AB; 5, TMZ plus 3-AB plus BG; 6, TMZ plus 6-AN; 7, TMZ plus 6-AN plus BG; 8, TMZ plus PD128763; 9, TMZ plus PD128763 plus BG.

Figure 10. The effect of MX plus TMZ on growth of SW480 xenograft.

Figure 11. The effect of MX plus TMZ on growth of HCT116 xenograft.

Figure 12A shows the effects of BCNU alone and BCNU in combination with MX on HCT116 tumors grown in nude mice. Figure 12B shows the effects of BCNU alone and BCNU in combination with MX on HCT116-Ch3 tumors grown in nude mice. Figure 12C shows the effects of BCNU alone and BCNU in combination with MX on SW480 tumors grown in nude mice.

Figure 13 shows the effects of hydrogen peroxide (H_2O_2) alone and H_2O_2 in combination with MX on HCT116 cells in a growth inhibition assay.

DEFINITIONS

To facilitate understanding of the invention, a number of terms are defined below.

As used herein "agent" or "drug" is used herein to denote a chemical compound, a mixture of chemical compounds, a biological macromolecule, or an

extract made from biological materials such as bacteria, plants, fungi, or animal (particularly mammalian) cells or tissues that are suspected of having therapeutic properties. The agent or drug may be purified, substantially purified or partially purified.

5 As used herein "agonist" refers to molecules or compounds which mimic the action of a "native" or "natural" compound. Agonists may be homologous to these natural compounds in respect to conformation, charge or other characteristics. Thus, agonists may be recognized by, *e.g.*, receptors expressed on cell surfaces. This recognition may result in physiologic and/or biochemical changes within the cell, such
10 that the cell reacts to the presence of the agonist in the same manner as if the natural compound was present.

As used herein "antagonist" refers to molecules or compounds which inhibit the action of a "native" or "natural" compound. Antagonists may or may not be homologous to these natural compounds in respect to conformation, charge or other
15 characteristics. Thus, antagonists may be recognized by the same or different receptors or molecules that are recognized by an agonist. Antagonists may have allosteric effects which prevent the action of an agonist (*e.g.*, by modifying a DNA adduct, or antagonists may prevent the function of the agonist (*e.g.*, by blocking a DNA repair molecule).

20 As used herein, the term "fusion protein" refers to a chimeric protein containing the protein of interest joined to an exogenous protein fragment. The fusion partner may provide a detectable moiety, may provide an affinity tag to allow purification of the recombinant fusion protein from the host cell, or both. If desired, the fusion protein may be removed from the protein of interest by a variety of enzymatic or
25 chemical means known to the art.

As used herein, the term "purified" or "to purify" refers to the removal of one or more contaminants from a sample. The present invention contemplates purified compositions (discussed above).

As used herein, the term "partially purified" refers to the removal of a moderate
30 portion of the contaminants of a sample to the extent that the substance of interest is

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recognizable by techniques known to those skilled in the art as accounting for a measurable amount of the mixture. Preferably, the compound of interest is at least 5% of the total preparation and up to 50% of the total preparation.

5 As used herein, the term "substantially purified" refers to the removal of a significant portion of the contaminants of a sample to the extent that the substance of interest is recognizable by techniques known to those skilled in the art as the most abundant substance in the mixture.

10 As used herein the term "portion" when in reference to a protein (as in "a portion of a given protein") refers to fragments of that protein. The fragments may range in size from four amino acid residues to the entire amino acid sequence minus one amino acid. In one embodiment, the present invention contemplates "functional portions" of a protein. Such portions are "functional" if they contain a binding region (i.e. a region having affinity for another molecule) and such binding can take place (i.e. the binding region functions, albeit with perhaps lower affinity than that observed for the full-length protein). Such "functional portions" of the gene product are 15 typically greater than 10 amino acids in length, and more typically greater than 50 amino acids in length, and even more typically greater than 100 amino acids in length. "Functional portions" may also be "conserved portions" of the protein. The alignment of the various gene products permit one skilled in the art to select conserved portions of the protein (i.e. those portions in common between two or more species) as well as 20 unconserved portions (i.e. those portions unique to two or more species). In one embodiment, the present invention contemplates conserved portions 10 amino acids in length or greater, and more typically greater than 50 amino acids in length.

25 "Staining" shall be defined as any number of processes known to those in the field that are used to better visualize, distinguish or identify a specific component(s) and/or feature(s) of a cell or cells.

30 "Immunofluorescence" is a staining technique used to identify, mark, label, visualize or make readily apparent by procedures known to those practiced in the art, where a ligand (usually an antibody) is bound to a receptor (usually an antigen) and such ligand, if an antibody, is conjugated to a fluorescent molecule, or the ligand is

then bound by an antibody specific for the ligand, and said antibody is conjugated to a fluorescent molecule, where said fluorescent molecule can be visualized with the appropriate instrument (*e.g.* a fluorescent microscope).

5 "Antibody" shall be defined as a glycoprotein produced by B cells that binds with high specificity to the agent (usually, but not always, a peptide), or a structurally similar agent, that generated its production. Antibodies may be produced by any of the known methodologies [Current Protocols in Immunology (1998) John Wiley and Sons, Inc., N.Y.] and may be either polyclonal or monoclonal.

10 "In operable combination", "in operable order" and "operably linked" as used herein refer to the linkage of nucleic acid sequences in such a manner that a nucleic acid molecule capable of directing the transcription of a given gene and/or the synthesis of a desired protein molecule is produced. The term also refers to the linkage of amino acid sequences in such a manner so that a functional protein is produced.

15 "Antigen" shall be defined as a protein, glycoprotein, lipoprotein, lipid or other substance that is reactive with an antibody specific for a portion of the molecule.

"Morphology" shall be defined as the visual appearance of a cell or organism when viewed with the eye, a light microscope, a confocal microscope or an electron-microscope, as appropriate.

20 "Patient" shall be defined as a human or other animal, such as farm animals or laboratory animals (*e.g.* guinea pig or mice) capable of having cell cycle (influenced) determined diseases, either naturally occurring or induced, including but not limited to cancer.

25 A "wild type" (wt) cell or cell line shall be defined as a cell or cell line that retains the characteristics normally associated with that type of cell or cell line for the physiological process or morphological characteristic that is being examined. It is permissible for the cell or cell line to have non-wild type characteristics for physiological process or morphological characteristics that are not being examined as long as they do not appreciably affect the process or characteristic being examined.

GENERAL DESCRIPTION OF THE INVENTION

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5 This invention generally relates to novel compositions and methods for the treatment of certain cancers. Additionally, this invention relates to novel compositions and methods to screen drugs useful for the treatment of certain cancers. In DNA repair competent cells, DNA adducts formed by methylating agents may be efficiently repaired or may be sites of both mutagenic and cytotoxic damage. In this process, the cellular response is specific for each of the DNA adducts formed. Perhaps the best studied is the response to O⁶-methylguanine (O⁶mG). This adduct may be repaired in a single step reaction by O⁶-alkylguanine-DNA alkyltransferase (AGT), however, 10 saturation of this protein by an excess of adducts or inhibition by O⁶-benzylguanine (BG) results in residual adducts which are both cytotoxic and mutagenic (Pegg *et al.* "Structure, function and inhibition of O⁶-alkylguanine-DNA AGT" *Prog. Nucleic Acid Res. Mol. Biol.* 51:167-233, 1995.). Cytotoxicity results from recognition of this adduct by components of the mismatch repair (MMR) system, a 5 or 6 protein 15 complex which recognizes O⁶mG : thymine base pairs formed by DNA replication past O⁶mG, and excises thymine and surrounding bases, resulting in DNA strand breaks. However, a thymine is preferentially reincorporated opposite the persisting O⁶mG, triggering MMR function again. It has been hypothesized that this repetitive aberrant repair process increases DNA double strand breaks and acts as a trigger of apoptosis 20 (Fink *et al.* "The role of DNA mismatch repair in drug resistance" *Clin. Cancer Res.* 4:1-6, 1998.).

MMR deficiency results in inability to process the O⁶mG:T mispair. Consequently cells replicate DNA past O⁶mG lesions without cell cycle arrest, chromosomal aberrations or apoptosis and, therefore, survive in the face of persistent 25 DNA damage (Branch *et al.* "Defective mismatch binding and a mutator phenotype in cells tolerant to DNA damage" *Nature* 362:652-654, 1993; Kat *et al.* "An alkylation-tolerant, mutator human cell line is deficient in strand-specific mismatch repair" *Proc. Natl. Acad. Sci. USA* 90:6424-6428, 1993; Karran and Bignami "DNA damage tolerance, mismatch repair and genome instability" *Bioessays* 16:833-839,

1994; Griffin *et al.* "DNA mismatch binding and incision at modified guanine bases by extracts of mammalian cells: implications for tolerance to DNA methylation damage" *Biochemistry* 33:4787-4793, 1994; Liu *et al.* "Mismatch repair mutations override alkyltransferase in conferring resistance to temozolomide by not to
5 1,3-bis(2-chloroethyl)nitrosourea" *Cancer Res.* 56:5375-5379, 1996). The presence of MMR deficiency in a number of colon cancer cell lines allowed the present inventors the opportunity to evaluate the relative contribution of this DNA repair defect in resistance to the methylating chemotherapeutic agent, temozolomide (TMZ), where we found that MMR deficiency resulted in 35-60 fold resistance to TMZ in cells defective
10 in either MLH1 or GTBP even after inhibition of AGT by BG (Liu *et al.* "Mismatch repair mutations override alkyltransferase in conferring resistance to temozolomide by not to 1,3-bis(2-chloroethyl)nitrosourea" *Cancer Res.* 56:5375-5379, 1996).

While O⁶mG is the best studied cytotoxic DNA adduct, it is not the most abundant. TMZ like other methylating agents, also forms N⁷-methylguanine (N⁷mG) and N³-methyladenine (N³mA) DNA adducts at frequencies 11 and 1.5 times that of O⁶mG (our unpublished results). These DNA adducts are efficiently removed by base excision repair (BER) and appear to contribute little to cytotoxicity. In the first step of BER, a series of glycosylases recognize abnormal bases such as N³mA and N⁷mG
15 (O'Connor *et al.* "Isolation and structure of a cDNA expressing a mammalian 3-methyladenine-DNA glycosylase" *EMBO J.* 9:3337-3342, 1990; Samson *et al.* "Cloning and characterization of a 3-methyladenine DNA glycosylase cDNA from human cells whose gene maps to chromosome 16" *Proc. Natl. Acad. Sci. USA* 88:9127-9131, 1991), the T:G mismatch (Neddermann *et al.* "Functional expression of soluble human interleukin-11 (IL-11) receptor alpha and stoichiometry of in vitro
20 IL-11 receptor complexes with gp130" *J. Biol. Chem.* 271:12767-12774, 1996), and deaminated bases such as hypoxanthine/oxidized 8-oxo-7,8-dihydroguanine or uracil:A (Vollberg *et al.* "Isolation and characterization of the human uracil DNA glycosylase gene" *Proc. Natl. Acad. Sci. USA* 86: 8693-8697, 1989; Olsen *et al.* "Molecular cloning of human uracil-DNA glycosylase, a highly conserved DNA repair enzyme"
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EMBO J. 8:3121-3125, 1989; Radicella *et al.* "Cloning and characterization of hOGG1, a human homolog of the OGG1 gene of *Saccharomyces cerevisiae*" *Proc. Natl. Acad. Sci. USA* 94:8010-8015, 1997; Rosenquist *et al.* "Cloning and characterization of a mammalian 8-oxoguanine DNA glycosylase" *Proc. Natl. Acad. Sci. USA* 94:7429-7434, 1997). Following enzymatic or spontaneous hydrolysis of the N-glycosidic bond and release of the abnormal base, AP (apurinic/aprimidinic) endonuclease hydrolyzes the phosphodiester backbone 5' to the lesion and dRpase (a DNA deoxyribophosphodiesterase and its activity is associated with polymerase β) excises the residual dRp, generating a gap of one nucleotide. DNA polymerase β fills the gap and DNA ligase seals the nick. This pathway is called short-patch BER. An alternative pathway for BER involves DNA synthesis to fill a gap of 2 to 13 nucleotides. This long patch repair requires proliferating cell nuclear antigen (PCNA) and PCNA-dependent DNA polymerase (Willson "Mammalian base excision repair and DNA polymerase beta" *Mutation Res.* 407:203-215, 1998).

Poly-(ADP-ribose)-polymerase (PARP) acts as a nick sensor of DNA strand breaks by itself or interaction with XRCC1 and involves in BER. PARP binds damaged DNA, resulting in autoribosylation. The modified protein then releases and allows other proteins to access and repair DNA strand breaks (Willson "Mammalian base excision repair and DNA polymerase beta" *Mutation Res.* 407:203-215, 1998; Molinete *et al.* "Over production of the poly (ADP-ribose) polymerase DNA-binding domain blocks alkylation-induced DNA repair synthesis in mammalian cells" *EMBO J.* 12:2109-2117, 1993; Caldecott *et al.* "XRCC1 polypeptide interacts with DNA polymerase β and possibly poly (ADP-ribose) polymerase, and DNA ligase III is a novel molecular 'nick-sensor' in vitro" *Nucleic Acids Res.* 24:4387-4394, 1996). Therefore, PARP participates in BER after nick formation in both short patch and long patch repair. It appears most active in the alternative (long patch repair) pathway for BER.

BER as a therapeutic target to increase the cytotoxicity of methylating agents has been documented. Cells deficient in DNA polymerase β or blocked in expression

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of AP endonuclease by antisense oligonucleotides are sensitized to methylating agents (Sobol *et al.* "Requirement of mammalian DNA polymerase- β in base-excision repair" *Nature* 379:183-186, 1996; Walker *et al.* "A role for the human DNA repair enzyme HAP1 in cellular protection against DNA damaging agents and hypoxic stress" *Nucleic Acids Res* 22:4884-4889, 1994). In addition, mice deficient in N³-methyladenine DNA glycosylase exhibited increased sensitivity to alkylating drugs like BCNU and mitomycin C (Engelward *et al.* "Repair-deficient 3-methyladenine DNA glycosylase homozygous mutant mouse cells have increased sensitivity to alkylation-induced chromosome damage and cell killing" *EMBO* 15:945-952, 1996). On the other hand, overexpression of the N³-methyladenine DNA glycosylase, which increases the number of AP sites formed, also increases the cytotoxicity of methylating agents (Coquerelle *et al.* "Overexpression of N-methylpurine-DNA glycosylase in Chinese hamster ovary cells renders them more sensitive to the production of chromosomal aberrations by methylating agents- a case of imbalanced DNA repair" *Mutation Res.* 336:9-17, 1995.). Finally, cells lacking PARP activity are more sensitive to alkylating agents, with increased apoptosis and chromosomal instability (Menissier-de Murcia *et al.* "Requirement of poly (ADP - ribose) polymerase in recovery from DNA damage in mice and in cells" *Proc. Natl. Acad. Sci. USA* 94:7303-7304, 1997; Boulton *et al.* "Potentiation of temozolomide-induced cytotoxicity: a comparative study of the biological effects of poly (ADP-ribose) polymerase inhibitors" *Br. J. Cancer*, 72:849-856, 1995). These data suggest that balanced expression of proteins in the base excision repair complex is important to the efficient processing of lesions. BER is an important mechanism of resistance to therapeutic methylating agents.

We examined two classes of agents which could inhibit the BER pathway to determine whether they would increase the cytotoxicity of methylating agents in colon cancer cells, particularly in cells deficient in MMR. Since MMR deficient cells are tolerant to O⁶mG formed by TMZ, any change in cytotoxicity observed after use of a BER inhibitor would be due to interruption in repair of N⁷mG and N³mA DNA adducts. Our first strategy was to combine methoxyamine (MX) with TMZ. MX has

been shown to react with the free aldehyde formed at the abasic site exposed by glycosylases and to reduce cleavage at AP sites in mammalian cells, suggesting that the MX bound abasic site is not a substrate for AP endonuclease (Fortini *et al.*

"Mutagenic processing of ethylation damage in mammalian cells: the use of methoxyamine to study apurinic/apyrimidinic site-induced mutagenesis" *Cancer Res.* 53:1149-1155, 1993). In the regard that AP sites modified by MX are relatively stable and must be converted to cytotoxic lesions, we hypothesized that MX would interrupt BER in cells and potentiate the cytotoxic effects of TMZ, even in MMR defective cells. The second strategy we used was to inhibit PARP with 3,4-dihydro-5-methoxyisoquinoline-1(2H)-one (PD128763), 3-aminobenzamide (3-AB) or 6-aminonicotinamide (6-AN) and to subsequently treat cells with TMZ. We hypothesize that inactivated PARP would affect short and long patch BER, destabilize strand breaks, reduce interaction with other proteins during repair of methylated DNA adducts and lead to cell death, again, in both MMR proficient and deficient cells.

As contemplated above, since MMR defective cell lines are remarkably resistant to methylating agents yet accumulate high levels of 3 methylating DNA adducts, O⁶mG, N³mA and N⁷mG, we reasoned that the interruption of repair of N³mA and N⁷mG adducts by the BER process would sensitize cells to methylating agents. To address this issue, we studied the effect of MX on potentiation of TMZ-cytotoxicity. MX interacts specifically with the tautomeric open-ring form of deoxyribose generated from the removal of an abnormal base by glycosylase. The MX modified AP site is relatively stable (Liuzzi and Talpeart-Borle "A new approach to the study of the base-excision repair pathway using methoxyamine" *J. Biol. Chem.* 260:5252-5258, 1985; Rosa *et al.* "Processing in vitro of an abasic site reacted with methoxyamine: a new assay for the detection of abasic sites formed in vivo" *Nucleic Acids Res.* 19: 5569-5574, 1991) and inhibits the cleavage of AP sites in DNA by AP endonuclease in mammalian cells. This has been shown to protect cells from cytotoxicity, mutagenicity induced by SN1-type ethylating agents, such as ENU but not SN2 alkylating agent, diethylsulfate and MMS (Fortini *et al.* "Mutagenic processing of

ethylation damage in mammalian cells: the use of methoxyamine to study
apurinic/aprimidinic site-induced mutagenesis" *Cancer Res.* 53: 1149-1155, 1993;
Fortini *et al.* "Methoxyamine modification of abasic sites protects CHO cells from the
cytotoxic and mutagenic effects of oxygen alkylation" *Carcinogenesis* 13: 87-93,
5 1992). Moreover, the protection was strictly time-dependent and was limited to the
short period (30 min) after exposure to the alkylating agents (Fortini *et al.*
"Methoxyamine modification of abasic sites protects CHO cells from the cytotoxic and
mutagenic effects of oxygen alkylation" *Carcinogenesis* 13: 87-93, 1992).

In our initial studies, we observed that MX reduced cleavage at AP sites and
10 decreased BER in human colon cancer cell extracts. However, we did not see
protection of these two cell lines from ENU cytotoxicity using longer duration
exposure to MX. The short duration of MX studied previously may not have the same
impact on BER inhibition as does a longer exposure to MX. Our results showed that
MX synergistically increased TMZ-induced cytotoxicity in human colon cancer cell
15 lines in both MMR proficient and deficient cells. A similar degree of enhanced
cytotoxicity was observed with MX and MMS and with TMZ as well. The effect of
BG inhibition of AGT was additive to the effect of MX only in the MMR proficient
SW480 cell line but not in the MMR defective HCT116 cell line. These data suggest
that O⁶mG DNA adducts do not contribute to the enhanced cytotoxic effect of TMZ by
20 MX. Furthermore, a similar degree of enhanced cytotoxicity was observed with MX
and MMS as with TMZ, again implicating N³mA and N⁷mG induced abasic sites as
the major targets for MX. In our recent studies, a prolonged exposure to low dose
MX results in even greater potentiation of TMZ cytotoxicity.

The mechanisms of MX-enhanced cytotoxicity of methylating agents in colon
25 cancer cells have not been fully understood. It is fair to suggest that MX enhanced the
cytotoxic effect of TMZ due to i). the MX-AP site complex is able to block the AP
endonucleolytic step of the BER pathway; ii). The persistence of abasic sites may
increase topoisomerase II mediated DNA cleavage (Kingma and Osheroff "Apurinic
sites are position-specific topoisomerase II poisons" *J. Biol. Chem.* 272: 1148-1155,

1997); iii). AP sites inhibit DNA replication and trigger programmed cell death (Robertson *et al.* "Down-regulation of apurinic/aprimidinic endonuclease expression is associated with the induction of apoptosis in differentiating myeloid leukemia cells" *Cell Growth & Differentiation* 8: 443 -449, 1997).

5 Under normal circumstances, TMZ produces strand breaks during BER mediated repair of N⁷mG and N³mA adducts which are efficiently repaired and do not contribute to cytotoxicity until high concentrations of adducts are achieved. When DNA strand breaks are present, one component of the response is recognition, binding and activation of PARP. Activated PARP leads to autoribosylation, and this in turn
10 facilitates access of repair enzymes to DNA damage (Willson "Mammalian base excision repair and DNA polymerase beta" *Mutation Res.* 407: 203-215, 1998; Buschfort *et al.* "DNA excision repair profiles of normal and leukemic human lymphocytes: functional analysis at the single-cell level" *Cancer Res.* 57: 651-658, 1997) and appears to enhance processing of stand breaks and religation by
15 polymerase-β and ligase I (Althaus *et al.* "Histone shuttle driven by the automodification cycle of poly(ADP-ribose)polymerase" *Environ Mol Mutagen*, 22: 278-282, 1993). In the alternative BER pathway, PARP interacts with XRCC1 to facilitate repair (Caldecott *et al.* "XRCC1 polypeptide interacts with DNA polymerase β and possibly poly (ADP-ribose) polymerase, and DNA ligase III is a novel molecular 'nick-sensor' in vitro" *Nucleic Acids Res.* 24: 4387-4394, 1996). It seems likely that
20 PARP plays an important role in communication between repair proteins and stability of the repair complex involved in BER. This suggests that inhibition of PARP leads to an impaired ability to rejoin DNA strand breaks which can initiate both apoptotic and non-apoptotic cell death cascades and thereby increase cytotoxicity of TMZ
25 (Wedge *et al.* "3-Aminobenzamide and /or O⁶benzylguanine evaluated as an adjuvant to temozolomide or BCNU treatment in cell lines of variable mismatch repair status and O⁶-alkylguanine-DNA alkyltransferase activity" *Br. J. Cancer* 74: 1030-1036, 1996). Our results support this hypothesis. Potentiation of cytotoxicity of methylating

agent with PARP inhibitors was observed with a marked increase in apoptosis and PARP cleavage.

Results of BCNU combined with either MX or PD128763 are in sharp contrast to TMZ; little if any potentiation is observed in the absence of BG in either cell line.

5 This suggests that while BER appears to process BCNU induced crosslinks, inhibition of BER in this manner has little impact on BCNU toxicity. One of the best studied BCNU-induced lethal lesions is the N³C-N¹G interstrand cross-link formed after initial chloroethyl monoadducts at O⁶G and cyclic rearrangement to N¹, O⁶-ethanoguanine (Wilson *et al.* "Life without DNA repair" *Proc. Natl. Acad. Sci. USA* 94: 7303-7304, 10 1997). However, treatment of cells with BCNU also produces alkylated bases that may be labile and spontaneously result in breakage or nicking of the phosphoribosyl backbone (Gonzaga *et al.* "Identification of the cross-link between human O⁶-methylguanine-DNA methyltransferase and chloroethylnitrosourea-treated DNA" *Cancer Res.* 52:6052 -6058, 1992). Since PARP has been shown to bind to 15 BCNU-induced DNA nicks *in vitro* (Prakash and Gibson "Sequence-selective depurination, DNA interstrand cross-linking and DNA strand break formation associated with alkylated DNA" *Carcinogenesis* 13: 425-431, 1992), it is reasonable to assume that PD128763 might increase BCNU-cytotoxicity. However, our studies showed only minor enhancement of toxicity in HCT116 cells and no enhancement in 20 SW480 cells. Even though methyladenine DNA glycosylase has been implicated in BCNU cross-link repair and its absence sensitizes cells to BCNU (Malapetsa *et al.* "Identification of a 116 kDa protein able to bind 1,3-bis (2-chloroethyl)-1-nitrosourea-damaged DNA as poly (ADP-ribose) polymerase" *Mutation Res.* 362: 41-50, 1996), we did not observe sensitization to BCNU by 25 treatment with MX in the absence of BG. In the presence of BG, MX potentiated BCNU toxicity, indicating that MX may interfere with DNA cross-link repair pathway and suggesting that BER may be involved in repair of the N¹O⁶ethanoguanine cross-link which is not formed if AGT reacts with the O⁶chloroethylguanine adduct. Taken together, these data suggest a different reaction of MX with damaged DNA

induced by BCNU compared to TMZ. With TMZ, MX-increased cytotoxicity is associated with AP sites generated from repair of N⁷mG and N³mA DNA adducts formed by methylating agent but with BCNU, it might be the O⁶ lesion induced cross-link which controls BCNU toxicity.

5 While an understanding of the precise mechanism is not needed to practice the present invention, it appears that the apoptosis mediates both MX and PD128763-enhanced cytotoxicity of TMZ. Increased apoptosis was observed in MMR wt SW480 cells but not in MMR deficient HCT116 cells after treatment with BG and TMZ. This suggests that MMR processing of O⁶mG is a potent apoptosis-inducing event (Matijasevic *et al.* "Protection against chloroethylnitrosourea cytotoxicity by eukaryotic 3-methyladenine DNA glycosylase" *Proc. Natl. Acad. Sci. USA*, 10 90:11855-11859, 1993). Although the biological and functional consequences of PARP and its cleavage in apoptosis still remain to be further identified, it has been demonstrated that PARP is rapidly and specifically cleaved during apoptosis (Kaina *et al.* "Chromosomal instability, reproductive cell death and apoptosis induced by 15 O⁶-methylguanine in Mex-, Mex+ and methylation-tolerant mismatch repair compromised cells: facts and models" *Mutation Res.* 381: 227-241, 1997; Lindahl *et al.* "Post-translational modification of poly(ADP-ribose) polymerase induced by DNA strand breaks" *Trends Biochem Sci* 20: 405-411, 1995). PARP cleavage was observed 20 in both SW480 and HCT116 cells after treatment with either MX or one of the PARP inhibitors and TMZ, confirming activation of apoptotic pathways.

We noted that arrest at cell cycle checkpoints paralleled the cellular response to DNA damage and that these were dependent on MMR and BER pathways. MMR wt SW480 cells were sensitive to TMZ alone with arrest in the S and G2 phases (Fink *et al.* "The role of DNA mismatch repair in drug resistance" *Clin. Cancer Res.* 4: 1-6, 25 1998). The S and G2 arrest were potentiated by MX or by PD128763 despite the fact that SW480 is a p53 mutant cell line. In contrast, even high levels of DNA adducts formed by TMZ in the MMR deficient HCT116 cells did not induce cell cycle checkpoint arrest despite the fact that p53 is wild type in this cell line. This

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dysregulation of damage-induced cell cycle checkpoint control appeared due to failure of processing O⁶mG lesions in MMR deficient cells. However after combined treatment with TMZ and either MX or PD128763, HCT116 cells showed S/G2 phase arrest and apoptosis. These results are consistent with previous studies of cell cycle changes following MMS exposure or other compounds that produce 90% N³mA (Lazebnik *et al.* "Cleavage of poly(ADP-ribose) polymerase by a proteinase with properties like ICE" *Nature* 371: 346-347, 1994; Morris *et al.* "Flow cytometric evaluation of cell-cycle progression in ethyl methanesulfonate and methyl methanesulfonate-exposed P3 cells: relationship to the induction of sister-chromatid exchanges and cellular toxicity" *Environ. Mol. Mutagen.* 18: 139-149, 1991) and the prolonged G2 arrest observed in PARP knockout mice or derived cell lines (Menissier-de Murcia *et al.* "Requirement of poly (ADP - ribose) polymerase in recovery from DNA damage in mice and in cells" *Proc. Natl. Acad. Sci. USA* 94: 7303-7304, 1997) following DNA damage. These data indicate that both SW480 and HCT116 cells have a similar response to persistent N⁷mG and N³mA lesions, following interruption of BER.

Again, while an understanding of the mechanism is not necessary to practice the invention, it appears that disrupted BER processing of non O⁶mG, most likely N⁷mG and N³mA DNA adducts formed by TMZ is cytotoxic to colon cancer cell lines. This may be particularly important in MMR deficient cells which are resistant to TMZ alone due to the failure to recognize O⁶mG DNA adducts. These studies provide evidence that disrupting repair of N⁷mG and N³mA by inhibiting BER or PARP may improve the therapeutic efficacy of methylating agents.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Generally, the nomenclature used hereafter and the laboratory procedures in cell culture, tissue culture, tumor biology, and molecular genetics described below are those well known and commonly employed in the art. Standard techniques are used for cell culture methods, experimental design and compound formulation and

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nomenclature. Generally chemical reactions and purification steps are performed according to the manufacturer's specifications. The techniques and procedures are generally performed according to conventional methods in the art and various general references (see, generally, Sambrook et al. Molecular Cloning: A Laboratory Manual, 2d ed. (1989) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., and Current Protocols in Molecular Biology (1996) John Wiley and Sons, Inc., N.Y., which are incorporated herein by reference) which are provided throughout this document. All the information contained therein is incorporated herein by reference.

Assays for detecting the ability of agents to inhibit or enhance TMZ-mediated tumor reduction provide for facile high-throughput screening of agent banks (e.g., compound libraries, peptide libraries, and the like) to identify potentiator. Such TMZ potentiator may be further developed as potential therapeutics and diagnostic or prognostic tools for diverse types of cancers. Likewise, TMZ and various potentiator may be useful in the treatment or diagnostics of other diseases. For example, such combinations of drugs may be directed to virally or parasitically infected cells

Thus, the present invention provides a method for potentiating the toxicity of temozolomide in human cancer cells by administering an APE or a PARP inhibiting amount of an APE or PARP inhibitor. The present invention also provides a product comprising temozolomide and an APE or PARP inhibitor as a combined preparation for simultaneous, separate or sequential administration in said treatment of human cancer cells. Additionally the present invention provides a method for the screening of compounds that may potentiate the toxicity of temozolomide by the mechanisms of inhibiting APE or PARP, or by other mechanisms.

I. Potentiation of Temozolomide Treatment for Cancer

The present invention contemplates compositions and methods for the potentiation of the elimination of cancer cells by temozolomide. While not limiting the invention to any particular mechanism, it is believed that temozolomide works as an anticancer agent by producing the DNA adducts O⁶mG, N⁷mG and N³mA. An accumulation of adduct results in the apoptotic elimination of the cell. However, the

O⁶mG adducts are typically repaired by the MMR system and the N⁷mG and N³mA adducts are typically repaired by the base excision repair (BER) system. After removal of the abnormal base by methylpurine glycosylase, AP endonuclease (APE) cleaves the back bone and removes the sugar, allowing DNA repair. If the number of adducts produced outstrips the cells ability to repair the damage, or the BER repair mechanism is rendered ineffective, damage will accumulate and the cell will be eliminated through an apoptotic mechanism. Methoxyamine reacts with the abasic site and prevents APE cleavage, disrupting DNA repair. Methoxyamine has been shown in the present invention to potentiate the activity of temozolomide in the treatment of cancer. PARP aids in the repair of DNA strand breaks induced during MMR and BER. As such, inhibitors of PARP activity have been shown in the present invention to be effective in potentiating the activity of temozolomide in the elimination of cancer cells. The effectiveness of temozolomide as an anticancer agent can be greatly enhanced when used in conjunction with a potentiator that prevents the repair of the DNA adducts created as a result of treatment of the cell with temozolomide.

Temozolomide has been found to be most preferably administered in repeat dosages on consecutive days, and the dramatic potentiation effects of the present invention are realized in the highly preferred regimen involving the administration of MX or a PARP inhibitor prior to, or concurrent with, the administration of each dose.

Preferably, this administration of an APE or PARP inhibitor is repeated over a period of several days and is prior to the administered doses of temozolomide. Repeat doses can be administered at 1, 2, 3, 4 or 5 days or more with the preferred period of therapy determined by the response of the tumor to the treatment.

In a preferred embodiment, the APE or PARP inhibitor is administered in an APE or PARP inhibiting amount, *i.e.* an amount sufficient to sensitize the tumor *in vivo* without causing undue sensitization of normal tissue, when the APE or PARP inhibitor is used concurrently with temozolomide.

The amount of the APE or PARP inhibitor used in the present invention varies according to the degree of the effective amount required for the treating of tumor cells. A suitable dosage is that which will result in a concentration of the APE or PARP

inhibitor in the tumor cells to be treated which results in the depletion of the APE or PARP activity, respectively. If desired, some tumor cells can be obtained by biopsy and can be tested *in vitro* for their sensitivity prior to *in vivo* treatment.

The neoplasms for which temolozomide is a particularly suitable treatment include carcinomas, melanomas, sarcomas, lymphomas and leukaemias, with specific utility for astrocytoma, gliomas, malignant melanoma, chronic lymphocytic leukaemia, lung cancer and best cancer.

Typical dosage ranges of temolozomide are generally between 0.1 and 200, preferably between 1 and 20 mg/kg body weight per day. The amount of APE inhibitor or PARP inhibitor necessary for the potentiation of temolozomide is dependent upon the amount of APE or PARP, respectively, normally present in the cancer cell type. A cancer cell having higher levels of APE or PARP will be potentiated more dramatically by the preadministration or simultaneous administration of the APE or PARP inhibitor. Additionally, the amount of APE or PARP inhibitor necessary depends on the age and condition of the patient, the severity of the cancer being treated and the particular inhibitor being utilized.

II. Drug Screens

As noted above, the present invention contemplates screening assays for identifying compounds that inhibit tumor growth. The present invention contemplates a screening assay utilizing cell lines that are resistant to temolozomide to screen for compounds that potentiate the anticancer properties of temolozomide. In one embodiment, a temolozomide resistant cell line is pretreated with the compound suspected of potentiating the anticancer properties of temolozomide followed by treatment with temozolomide. Cell growth and/or apoptosis are then measured. In another embodiment, the compound suspected of potentiating the anticancer temozolomide is given simultaneously with temozolomide.

Additionally, the present invention contemplates the screening of compounds by the use of xenographs. In one embodiment, tumor cells are injected into immune compromised mice (or other suitable animal) and allowed to grow. Mice carrying the

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tumors are then treated with the compound suspected of potentiating temozolomide in parallel with or prior to treatment with temozolomide. Tumor size is then measured to follow the effect of the treatment.

5 It is not intended that the present invention be limited by the nature of the drugs screened in the screening assay of the present invention. A variety of compounds, including peptides, organic compounds, nonorganic compounds, as well as, formulations of more than one compound, are contemplated.

10 It is also not intended that the present invention be limited by the particular tumor cells used for the drug testing. A variety of tumor cells (for both positive and negative controls) are contemplated (including but not limited to cells set forth above and in the examples below).

15 It is also not intended that the present invention be limited by the mechanism by which the compound suspected of potentiating TMZ works. For example, the compound may work by inhibiting APE cleavage, by inhibiting other DNA repair mechanisms (*e.g.* PARP) or by other as of yet unidentified mechanisms.

20 It is contemplated that the invention be utilized in the screening of compounds by high-throughput screening methods. For example, automated systems and microscale assay systems are contemplated. Furthermore, it is contemplated that the invention can be used for the screening of compound libraries (*e.g.* drug libraries) utilizing these high throughput screening methods.

EXPERIMENTAL

Materials and Methods

25 **Chemicals and Reagents.** BG was generously provided by Dr. Robert Moschel (Frederick Cancer Research and Development Center, National Cancer Institute). Stock solution was made in dimethylsulfoxide. TMZ and BCNU were obtained from the Drug Synthesis and Chemistry Branch, Drug Therapeutic Program, National Cancer Center Institute. PD128763 was a gift from Park-Davis Pharmaceutical Division. 6-AN, 3-AB, MX and methylmethane sulfonate (MMS) were purchased from Sigma (ST. Louis, MO). Stock solutions of PD128763, 3-AB and 6-AN were

prepared by dissolving in dimethylsulphoxide and added to cell culture at a final concentration of <1% DMSO, when cells were treated with these compounds. MX was dissolved in sterilized water (PH 7.0). All stock solutions were kept in -20°C. BCNU was prepared fresh in 0.5ml of 100% ethanol, diluted in PBS, and used with in 10 min.

Colony Survival Assay. SW480 cells were obtained from the ATCC. HCT116 cells were obtained from R. Boland, University of Michigan Medical Center. All cell lines were cultured in appropriate growth medium.

Cells (2000/dish) were plated, adhered for 18 hrs and treated with TMZ or MMS plus or minus modifiers such as BX, MX, 6-AN, 3-AB or PD128763, according to experimental protocol. After treatment, cells were washed and fresh medium was replaced. The cells were grown for a further 7 days prior to staining with methylene blue for determination of colonies containing more than 50 cells. Comparisons of drug induced cytotoxicity consisted of a calculation of the dose modification factor (DMF), defined as the ratio of the IC_{50} of either TMZ or MMS in the absence to that in the presence of indicted modifier(s), *i.e.* $DMF = IC_{50} \text{ for TMZ alone} / IC_{50} \text{ for TMZ plus modifier(s)}$.

Median Effect Analysis (MEA). MEA was used to determine the dose-response interactions between TMZ and either MX or PD128763. Drugs were combined at the ratio of the IC_{50} values for either TMZ and MX or TMZ and PD128763 as determined by survival / concentration curves. The combination was compared to the cytotoxicity of each drug alone in every experiment. The combination index (CI) was determined from colony-forming assays at increasing levels of cell killing, using analysis of multiple drug interaction program (Biosoft, Cambridge, United Kingdom) developed based on method of Chou and Talalay (Chou and Talalay "Quantitative analysis of dose-effect relationship: the combined effects of multiple drugs on enzyme inhibitors" In: G. Weber (ed.) *Advances in Enzyme Regulation*, pp. 27-55, New York: Pergamon

Press, 1983). CI values of less than or greater than 1 indicate synergy and antagonism, respectively., whereas a CI value of 1 indicates additivity of the drugs.

Flow Cytometry for Cell Cycle Distribution Analysis. For cell cycle analysis, 10^6 cells were plated in 100-mm tissue culture dishes and exposed to MX (6mM)/PD128763 (100 μ M) or MX (6mM)/PD128763 (100 μ M) plus TMZ (300 μ M) at 37°C. After 24h-72h of culture, cells were fixed in 80% ethanol and DNA was stained with 20 μ g/ml propidium iodide. DNA fluorescence of PI-stained cells was measured with an Elite ESP flow cytometer/cell sorter (Coulter, Miami, FL). Cell cycle distribution was analyzed with Modfit 5.2 program (Verity software, Topsham, MA) at least 10,000 cells per data point.

Western Blotting for PARP Cleavage Detection. Cell extracts were resolved by SDS-PAGE (12% polyacrylamide) in a Bio-Rad minigel apparatus at 150 V for 1 hr. Proteins were transferred onto PVDV membranes, using a Bio-Rad mini Trans-Blot cell for 1 hr at 100 V. The blotted membranes were blocked with 5% dry milk in TBS buffer and then probed for 2 hr with anti-PARP antibody C2-10 (Trevigen, Gaithersburg, MD). After three 5 min washes with TBS-Tween20 (0.05%), the blots were incubated with secondary antibody, anti-mouse HRPO-anti IgG for 1 hr (Amersham Life Science, Arlington Height IL). Antibody binding was visualized by ECL according to manufacturer's instructions (Amersham Life Science, Arlington Heights, IL).

Tumors in Nude Mice. Tumor cells (5×10^6) were injected into flanks of female athymic HSD nude mice, at 6-8 weeks of age. The tumors were measured with calipers using the National Cancer Institute formula: $V = L \text{ (mm)} \times I^2 \text{ (mm)} / 2$ where L is the largest diameter and I is the smallest diameter of the tumor. When the volume of the nodules has achieved about 100-150 mm³, tumor-bearing mice were assigned randomly for the control or treatment groups (3-5 mice / group).

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Antitumor Effects of TMZ and MX plus TMZ. Nude mice carrying tumors were received TMZ, MX or MX combined with TMZ daily for a period of 5 days of treatment. Doses used per group were as follows: a) TMZ alone, 120 mg/kg; b) MX alone, 2 mg/kg; c) BG plus TMZ, 30 mg/kg BG + 120 mg/kg; d) MX plus TMZ, 0.2 mg/kg MX + 120 mg/kg TMZ.

Experimental End Point. Tumor measurements were taken every 3 days. The relative tumor volume (V/V_0) was calculated by dividing the measured tumor volume (V) by the initial tumor volume (V_0) at day 0. Tumor responses were quantified by tumor regrowth delay. Tumor growth delays were calculated according to: tumor growth delay = $T_{2X} - C_{2X}$, where T_{2X} and C_{2X} represent the number of days treated and control tumors take to double in size from the day of treatment, respectively.

Toxicity Evaluation. Toxicity after treatment was evaluated by body weight measurements and peripheral WBC counting. Body weight was measured three times weekly from the first treatment until 2 weeks after the end of treatment. The weight loss was expresses as a percentage of the initial weight (initial weight - lowest weight / initial weight x 100%). Peripheral WBCs were monitored 5 days after the end of treatment compared with WBCs of control mice.

EXAMPLE I

Methoxyamine Potentiates Cytotoxicity of TMZ

We have previously reported the comparative cytotoxicity of TMZ and BG in the SW480 and HCT116 cells lines (Liu *et al.* "Mismatch repair mutations override alkyltransferase in conferring resistance to temozolomide by not to 1,3-bis(2-chloroethyl)nitrosourea" *Cancer Res.* 56: 5375-5379, 1996). In order to test whether MX would alter TMZ cytotoxicity, we treated SW480 and HCT116 with 6 mM MX (itself a nontoxic concentration) plus TMZ (0-1500 μ M) for 2 hr, with or without BG to abolish AGT mediated removal of O⁶mG DNA adducts. SW480 cells were moderately resistant to TMZ, with an IC₅₀ of 395 μ M which was reduced 14 fold

to 28 μ M by BG pretreatment. Greater resistance to TMZ was observed in MH1 defective cells, even after inhibition of AGT by BG (IC₅₀, 950 μ M). In both cell lines, MX potentiated the cytotoxic effect of TMZ with a DMF of 2.4 in SW480 and 3.1 in HCT116 (Fig. 1). In SW480 cells, additive effects of MX and BG were noted (IC₅₀ was reduced from 395 μ M to 6 μ M), whereas, with HCT116 cells no effect of BG was seen in the presence of MX (Fig. 1: A. HCT116 cells; B. SW480 cells. (■), TMZ alone; (◆), MX plus TMZ; (●), BG plus TMZ; (▲), BG and MX plus TMZ).

To further decipher the role of N³mA and N⁷mG DNA adducts in the relative absence of O⁶mG, we evaluated the effect of MX on MMS mediated cytotoxicity.

MMS is a methylating agent which produces far fewer O⁶mG adducts (0.3%) and a greater proportion of N³mA (10%) and N⁷mG adducts (87%) than TMZ (Chou and Talalay "Quantitative analysis of dose-effect relationship: the combined effects of multiple drugs on enzyme inhibitors" In: G. Weber (ed.) *Advances in Enzyme Regulation*, pp. 27-55, New York: Pergamon Press, 1983). The IC₅₀ of MMS was 0.82mM in SW480 and 1.4mM in HCT116 cells. This difference is smaller than the difference in the TMZ IC₅₀ between these cell lines, probably because the low level of O⁶mG adducts formed by MMS increases the impact of other DNA adducts. After cells were treated with MMS (0-3mM) plus 6mM MX for 1 hr, the IC₅₀ DMFs, compared to MMS alone, were 2.0 in SW480 and 2.3 in HCT116 (Fig. 2: A. HCT116 cells; B. SW480 cells. (■), MMS alone; (◆), MX plus MMS; (●), BG plus MMS; (▲), BG and MX plus MMS). These DMFs were similar to that observed with TMZ/ Compared to treatment of SW480 with BG plus TMZ (DMF was 14), BG plus MMS induced less enhancement of cytotoxicity (DMF was 6). When MMS was combined with BG and MX, >10 fold potentiation of cytotoxicity was observed SW480, whereas, no additive effect was seen in HCT116 cells. In SW480 cells, the increased MMS-toxicity by BG suggests that even the small number of O⁶mG adducts, they contribute to cytotoxicity. In HCT116 cells, increased cytotoxicity was only observed in the combination of TMZ with MX but not with BG, indicating that MX interfered with processing of N⁷mG and N³mA lesions which killed cells. From these data, we infer that MX had equal ability to interrupt BER in these two cell lines.

EXAMPLE 2

Inhibitors of PARP Modulate the Sensitivity of Cells to TMZ

Since inhibitors of PARP may interrupt BER and increase sensitivity to methylating agents, we examined whether inhibitors of PARP sensitize cells to TMZ. Fig. 3a and fig. 4a ((■), TMZ alone; (◆), BG plus TMZ; (●), PD128763 plus TMZ; (▲), BG and TMZ plus PD128763) display survival after combined treatment of TMZ with PD128763, 3-AB or 6-AN in both SW480 and HCT116 cells. In the SW480 cell line, PD128763 sensitized cells to TMZ with a DMF of 3.3 fold. The combination of PD128763, BG TMZ was even more toxic, with a DMF compared to TMZ alone of 36-fold. In HCT116 cells, the DMF for PD128763 and TMZ compared to TMZ alone was 5.0. However, the combination of PD128763, BG and TMZ had no greater effect than PD128763 and TMZ, indicating that persistent O⁶mG had no effect on cytotoxicity in this MMR defective cell line. Potentiation of TMZ cytotoxicity was also observed in both cell lines treated with two other PARP inhibitors, 3-AB (Fig. 3b, 4b: (■), TMZ alone; (◆), BG plus TMZ; (●), 6-AN plus TMZ; (▲), BG and TMZ plus 6-AN) and 6-AN (Fig. 4c, 4c: (■), TMZ alone; (◆), BG plus TMZ; (●), 3-AB plus TMZ; (▲), BG and TMZ plus 3-AB). Although the specific activity of these agents varied considerably, the DMFs were similar, 3-4-fold, for both 3-AB and 6-AN.

EXAMPLE 3

Synergistic Interaction Between TMZ and MX or PD128763

We investigated the nature of the reaction between TMZ and MX in the TMZ-resistant cell line, HCT116. These cells were incubated in the presence of a range of concentrations of TMZ (75-750μM), of MX (1.5-15.0mM), or constant molar ratio mixture of TMZ and MX (1:20) for 2 hr. HCT116 cells were also exposed to TMZ and PD128763 (62.5-625μM) alone and to a combination of (1:0.83) for 2 hr to analyze synergism. As shown in Fig. 5A and 5B, synergistic interaction ($CI < 1$, $p < 0.001$) was found in both SW480 cells and HCT116 cells using the combination of TMZ with either MX or PD128763. These results also showed a marked synergism at high Fa values. Since MDF for these combinations was similar to the observed with

SW480 cells, we conclude that BER inhibitor synergizes methylating agent cytotoxicity in both MMR deficient and proficient colon cancer cells.

EXAMPLE 4

Effect of BER Inhibitors on BCNU Cytotoxicity

To test whether MX is also able to sensitize colon cancer cells to chloroethylating agents, these two cell lines were pretreated with 6mM MX for 2 hr followed by BCNU. No enhancement of BCNU cytotoxicity by MX was observed (Fig. 6); the BCNU IC₅₀ was 45μM in HCT116 cells (Fig. 6A: (■), BCNU alone; (◆), MX plus BCNU; (●), BG plus BCNU; (▲), BG plus MX plus BCNU) and 27-29μM in SW480 cells, respectively, treated with BCNU alone or BCNU plus MX (Fig 6B: (■), BCNU alone; (◆), MX plus BCNU; (●), BG plus BCNU; (▲), BG plus MX plus BCNU). A greater sensitization to BCNU was observed in these two cell lines when cells were treated with MX plus BG and BCNU, in which the BCNU IC₅₀ for both cell lines was 5μM. However, most of the effect was potentiated due to BG, which increased BCNU cytotoxicity by 3-4 fold. As shown in Fig. 7, no sensitization to BCNU cytotoxicity was seen after treatment with PD128763 or BG and PD128763 ((■), BNCU alone; (◆), PD128763 plus BCNU; (●), BG plus BCNU; (▲), BG and PD128763 plus BCNU; (▶), BG, PD128763 and MX plus BCNU).

EXAMPLE 5

Effect of Inhibitors of BER on Cell Cycle Distribution and PARP Cleavage

The cell cycle and apoptosis response of SW480 and HCT116 cells was examined at various times after treatment with TMZ (300μM) alone or with either MX (6mM), PD128763 (100μM) or BG (25μM). After treatment, cells were divided into two aliquotes for analysis of cell cycle / apoptosis on days 1 and 3, and (see below) for detection of PARP cleavage. Cell cycle distribution was measured by flow cytometry according to DNA content and estimation of the duration of G1, S and G2/M was based on untreated, exponentially growing, asynchronous cells. MX and PD128763 alone did not affect the distribution of the cell cycle in these two cell lines

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(data not shown). At 24 hr, 75 - 90% of SW480 cells accumulated in S and G2 after treatment with TMZ alone and this G2/S phase arrest was more pronounced in cells pretreated with either MX or PD128763 (Fig. 8A). S/G2 arrest was still present at 3 days in cells treated with the combination of MX or PD128763 and TMZ (in both instances, 13 - 20% of the cells were apoptotic). In SW480 cells treated with TMZ alone, the G2/S block was less obvious at day 3 with only 8% of the cells showing evidence of apoptosis. In contrast, HCT116 cells had a normal cell cycle distribution after treatment with TMZ alone and no effect was seen with BG and TMZ. However, accumulation in S phase was observed (Fig 8B) 24 hr after treatment with PD128763 plus TMZ. At 72 hr, HCT116 cells had moved through S phase and thereafter, a significant portion of cells (90%) remained arrested in G2 with apoptosis present in 1% of the cells. A similar but less striking result was observed with MX and TMZ in HCT116 cells. By 72 hr, 60% of cells were still arrested in S and G2 and 10% of the cells were apoptotic.

Finally, as a marker of apoptosis induced cell death, we examined PARP cleavage after cells were treated with these drug combinations at 3 days (Fig 9). PARP cleavage was observed in SW480 cells after exposure to TMZ alone and TMZ plus BG, but was not seen in HCT116 cells with the same treatment, indicating that the apoptotic process is triggered when O⁶mG lesions are repaired by the MMR system. However, PARP cleavage was detected in MMR wild type and deficient cells treated with TMZ plus either MX or PARP inhibitors.

EXAMPLE 6

Acute Toxicity of MX

Three mice per group were injected (i.p.) with MX at doses of 1, 2, 4, 6, 10 and 15 mg/kg for the test of acute toxicity. MX was lethal at 15 mg/kg causing death within 10 mins. At 10 mg/kg, mice appeared to have decreased motility but recovered one hour after treatment. Lower doses did not appear toxic.

EXAMPLE 7

Antitumor Effect of TMZ or TMZ combined with either BG or MX on SW480 Tumor Xenograft

The response of SW480 xenograft to TMZ alone and combined treatments was shown in Fig 10 and Table 1. At the highest tolerable does of TMZ, 120 mg/kg, a tumor growth delay of approximately 12 days was noted, indicating that SW480 is sensitive to TMZ. When given 30 mg/kg of BG for one hour prior to 40 mg/kg of TMZ, tumor growth delay was enhanced by BG up to 3-fold. Slightly greater efficacy was seen by combining the BG with high doses of TMZ, but significant weight loss (maximum body weight loss from 26 to 20 g, 23%) and very low leukocyte counts (90% decreased) at day 5 after the last treatment were observed in the mice. In contrast ,mice treated with 0.2 mg/kg MX plus 120 mg/kg TMZ had an immediate cessation of tumor growth for 20 days and very slow regrowth of the tumor with tumor growth delays over 40 days ($p < 0.02$). There was no evidence of toxicity to mice with this combined treatment.

EXAMPLE 8

Antitumor Effect of TMZ or TMZ Combined with MX on HCT116 Tumor Xenograft

Mice bearing HCT116 xenograft appeared to be relatively sensitive to all treatment. Body weight loss was observed in all treated groups (Table 2). BG plus TMZ caused 28% body weight loss and toxic death (2 of 5 mice) at 5 days after treatment. As shown in Fig. 11, HCT116 xenografts treated either with TMZ or MX alone had similar growth rate and growth delays of tumor treated with 0.5 mg/kg MX plus 120 mg/kg TMZ were significantly prolonged compared with control tumors ($p < 0.05$).

EXAMPLE 9

While an understanding of the underlying mechanism is not necessary to the practice of the invention, and with the understanding that the invention is not to be

limited to any particular mechanism, the Inventors have shown that MX potentiates the antitumor effect of methylating agents such as temozolomide (TMZ) both *in vitro* and in human tumor xenograft models. A nonlimiting hypothesis for this potentiation is that the potentiation is mediated through the known ability of MX to bind

5 apurinic/pyrimidinic (AP) sites in DNA. Thus, in this model, one test of which is provided in the instant example, MX would be hypothesized to be useful in chemotherapy against cancer, and to enhance the therapeutic efficacy of a broad class of agents that produce damaged DNA to generate AP sites, thereby interrupting AP-

endonuclease (APE) in the base excision repair pathway.

10 This example shows that the combination of methoxyamine (MX) and 1,3-bis (chloroethyl) 2-nitrosourea (BCNU, a crosslinking agent known to produce AP sites) is effective in the treatment of cancer, using human tumor xenografts grown in nude mice.

Tumor cells (5×10^6) were injected into both flanks of female athymic nude mice, at 6-8 weeks of age. The tumors were measured with calipers using the National Cancer Institute formula: $V = L \text{ (mm)} \times I^2 \text{ (mm)} / 2$, where L is the largest diameter and I is the smallest diameter of the tumor. When the volume of tumor nodules has achieved about 100-150 mm³, tumor-bearing mice were assigned randomly for the control or the treatment groups (3-5 mice/group, 2 tumors/mouse).

15

20 Nude mice carrying tumor received a single injection (i.p.) of BCNU (30 mg/kg) or MX (2 mg/kg) combined with BCNU.

Tumor measurements are taken every 3 days. The relative tumor volume (V/V_0) was calculated by dividing the measured tumor volume (V) by the initial tumor volume (V_0) at day 0. Tumor responses were quantified by tumor regrowth delay.

25 Tumor growth delays were calculated according to the following formula: tumor growth delay = $T_{2x} - C_{2x}$, where T_{2x} and C_{2x} represent the number of days treated and control tumors take to double in size from the day of treatment, respectively.

30 The human colon cancer cell lines HCT116 (hMLH mut., p53 wt and AGT expressing) and HCT116-Ch3 (having restored MLH1 activity) were inoculated into

nude mice at 6-8 weeks of age. At the tumor sizes indicated above, the mice received a single i.p. injection, also as described above.

The effects on HCT116 tumors are depicted in Figure 12A. The effects on HCT116-Ch3 are depicted in Figure 12B. At a dose of 30 mg/kg, BCNU alone had a very mild effect on tumor growth. Combined administration of MX and BCNU produced significant tumor growth inhibition. Tumor growth delays (T2x-C2x) were 14-16 days in HCT116 tumors (Figure 12A) and HCT116-Ch3 (Figure 12B) respectively. A similar result was also observed in SW480 tumor (MMR wt, p53 mutant and AGT expressing). Tumor growth delays were 25 days for the SW480 tumors (Figure 12C).

Importantly, no systemic toxicity was noted in this combined treatment. In contrast, BCNU (at a dose of 25 mg/kg) combined with O6-benzylguanine (30 mg/kg), an AGT inhibitor being tested in clinical trials, caused toxic death at a 100% rate.

In *in vitro* studies, the inventors observed that cells exposed to MX combined with BCNU had increased DNA strand breaks [both single strand breaks (SSB) and double strand breaks (DSB) measured by comet assay and pulsed field gel] compared to BCNU alone in these two colon cell lines.

The potentiation of the BCNU-antitumor effect by MX suggests that the impact of the base excision repair pathway is important for BCNU-induced cytotoxicity and that MX adducted AP sites efficiently interrupt BER leading to cell death. Again, this is only a hypothesis, and the invention is not to be limited to any particular mechanism, and the practice of the invention does not require any knowledge of the underlying mechanism.

EXAMPLE 10

This example shows the combination of MX and an oxidizing agent for the treatment of cancer.

Again, while the present invention is not limited to any mechanism, BER comprises a ubiquitous series of biochemical pathways for the removal of oxidative damage to the nitrogenous bases in DNA. Initiating proteins of the BER are DNA

glycosylases, which hydrolyze the N-glycosylic bond between the deoxyribose sugar moiety and the DNA base, generating AP sites. AP sites can also be generated by spontaneous destabilization of N-glycosyl bonds, particularly after oxidative damage to the bases. The AP sites are the target for MX, resulting in the disruption of BER.

5 In this experiment, a growth inhibition assay was carried out with H_2O_2 , an oxidizing agent that mimics the effect of chemotherapeutic agents and radiation. Cells in the exponential phase were plated (1×10^4) in 24-well tissue culture plates. After attachment overnight, the cells were incubated with hydrogen peroxide (H_2O_2) (0-150 μ M) or MX (6 mM) plus H_2O_2 for 1 hr. Five days after treatment, the media was
10 discarded, and 100 μ l of MTT (5 mg/ml in PBS) was added in 1 ml fresh media to each well for 4 hrs at 37°C. The MTT solution was then removed, and the formazan crystals were dissolved in 200 μ l/well of DMSO for 20 min and changes in absorbance were determined at A_{540nm} . Percentage growth inhibition was calculated by comparison of the A_{540nm} reading from treated versus control cells. Drug concentrations that
15 resulted in and IC50 were determined from the plots of percentage growth inhibition versus the logarithm of the drug concentration.

As shown in Figure 13, MX results in significantly increased sensitivity of colon cancer cells, HCT116, to killing induced by H_2O_2 .

This result indicates that MX potentially increases the therapeutic efficacy of a
20 class of anticancer drugs that exert cytotoxicity mediated by oxidative damage in DNA, including but not limited to bleomycin, adriamycin and gamma radiation.

EXAMPLE 11

This example is directed to combinations of MX with members of a group of agents including, but not limited to hypoxanthine, 5-FU, uracil, IUdR, bleomycin,
25 adriamycin and gamma radiation.

The agents contemplated for combinations with MX in this example are agents which incorporate into DNA and are repaired by BER to form AP sites. Cell-growth inhibition was carried out with HCT116 cells to test MX-potentiated cytotoxicity of iododeoxyuridine (IUdR), which has been recognized as a radiosensitizing agent since

the early 1960's. As shown in Table 3, MX increases the killing effect of IUdR (or MX decreases the IC₅₀ of >>60 μ M for IUdR alone to 35 μ M) in MMR deficient HCT116 cells. This suggests that MX and IUdR combined with radiation therapy will further increase the radiosensitivity of tumor cells.

5 These experiments demonstrate that a significant enhancement of antitumor effect of TMZ by MX or PARP inhibitors in human colon cancer xenographs with mismatch repair proficiency and deficiency. Therefore, from the above it should be clear that the present invention provides a wide variety of ways to 1) screen for compounds that can potentiate TMZ treatment of cancer, 2) provide model systems for
10 the study of cancer treatment by agents that modulate DNA repair mechanisms and, 3) provide treatments for various cancers.

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Table 1. SW480 xenograft toxicity and tumor response

Group	Doses (mg/kg)	Toxicity (max. BW loss)	Tumor growth delays (Days)
TMZ	120	11% (27±1.9 - 24±2.4 g)	9.3 ± 1.2
MX	2	0 %	2.0 ± 0.5
BG+TMZ	30/120	23% (26±1.5 - 20±0.5 g)	10.3 ± 2.4
MX+TMZ	2/120	8 % (26±1.5 - 24±1.5 g)	27.0 ± 1.2

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Table 2. HCT116 xenograft toxicity and tumor response

Group	Doses (mg/kg)	Toxicity (max. BW loss)	Tumor growth delays (Days)
TMZ	120	20.8% (24±1.3 - 19±1.7 g)	2.5 ± 0.8
MX	2	8.3% (24±1.6 - 22±1.4 g)	2.2 ± 0.4
BG+TMZ	30/120	28.0% (25±0.8 - 18±0.5 g)*	NA
MX+TMZ	2/120	19.2% (26±1.3 - 21±0.4 g)	17.0 ± 1.2

* toxic death: 2/5 mice
NA, did not analyze

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TABLE 3

	IUdR	IUdR + MX (6 mM)
IC 50 (μ M)	>>60	35

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